

A Joint Graph Inference Case Study: the *C.elegans* Chemical and Electrical Connectomes

Li Chen^{*} Joshua T. Vogelstein[†] Vince Lyzinski[‡] Carey E. Priebe[§]

August 6, 2015

Abstract

We investigate joint graph inference for the chemical and electrical connectomes of the *Caenorhabditis elegans* roundworm. The *C.elegans* connectomes consist of 253 non-isolated neurons with known functional attributes, and there are two types of synaptic connectomes, resulting in a pair of graphs. We formulate our joint graph inference from the perspectives of seeded graph matching and joint vertex classification. Our results suggest that connectomic inference should proceed in the joint space of the two connectomes, which has significant neuroscientific implications.

The *Caenorhabditis elegans* (*C.elegans*) is a non-parasitic, transparent roundworm approximately one millimeter in length. The majority of *C.elegans* are female hermaphrodites. Maupas [1901] first described the worm in 1900 and named it *Rhabditis elegans*. It was later categorized under the subgenus *Caenorhabditis* by Osche [1952], and then, in 1955, raised to the generic status by Ellsworth Dougherty, to whom much of the recognition for choosing *C.elegans* as a model system in genetics is attributed [Riddle et al., 1997]. The long name of this nematode mixes Greek and Latin, where *Caeno* means recent, *rhabditis* means rod-like, and *elegans* means elegant.

Research on *C.elegans* rose to prominence after the nematode was adopted as a model organism: an easy-to-maintain non-human species widely studied, so that discoveries on this model organism might offer insights for the functionality of other organisms. The discoveries of caspases [Yuan et al., 1993], RNA interference [Fire et al., 1998], and microRNAs [Lee et al., 1993] are among some of the notable research using *C.elegans*.

^{*}Department of Applied Mathematics and Statistics, Johns Hopkins University, Baltimore, MD 21218, USA
corresponding author: lchen87@jhu.edu

[†]Department of Biomedical Engineering and Institute for Computational Medicine, Johns Hopkins University, Baltimore, MD 21218, USA

[‡]Johns Hopkins University Human Language Technology Center of Excellence, Baltimore, MD, 21218, USA

[§]Department of Applied Mathematics and Statistics, Johns Hopkins University, Baltimore, MD 21218, USA

Connectomes, the mapping of neural connections within the nervous system of an organism, provide a comprehensive structural characterization of the neural network architecture, and represent an essential foundation for basic neurobiological research. Applications based on the discovery of the connectome patterns and the identification of neurons based on their connectivity structure give rise to significant challenges and promise important impact on neurobiology. Recently there has been an increasing interest in the network properties of *C.elegans* connectomes. The hermaphrodite *C.elegans* is the only organism with a fully constructed connectome [Sulston et al., 1983], and has one of the most highly studied nervous systems.

Studies on the *C.elegans* connectomes traditionally focus on utilizing one single connectome alone [Varshney et al., 2011, Pavlovic et al., 2014, Sulston et al., 1983, Towlson et al., 2013], although there are many connectomes available. Notably, Varshney et al. [2011] discovered structural properties of the *C.elegans* connectomes via analyzing the connectomes’ graph statistics. Pavlovic et al. [2014] estimated the community structure of the connectomes, and their findings are compatible with known biological information on the *C.elegans* nervous system.

Our new statistical approach of joint graph inference looks instead at jointly utilizing the paired chemical and electrical connectomes of the hermaphrodite *C.elegans*. We formulate our inference framework from the perspectives of seeded graph matching and joint vertex classification, which we will explain in Section 2. This framework gives a way to examine the structural similarity preserved across multiple connectomes within species, and make quantitative comparisons between joint connectome analysis and single connectome analysis. We found that the optimal inference for the information-processing properties of the connectome should proceed in the joint space of the *C.elegans* connectomes, and using the joint connectomes predicts neuron attributes more accurately than using either connectome alone.

1 The Hermaphrodite *C.elegans* Connectomes

The hermaphrodite *C.elegans* connectomes consist of 302 labeled neurons for each organism. The *C.elegans* somatic nervous system has 279 neurons connecting to each other across synapses. There are many possible classifications of synaptic types. Here we consider two types of synaptic connections among these neurons: chemical synapses and electrical junction potentials. These two types of connectivity result in two synaptic connectomes consisting of the same set of neurons.

We represent the connectomes as graphs. A graph is a representation of a collection of interacting objects. The objects are referred to as nodes or vertices. The interactions are referred to edges or links. In a connectome, the vertices represent neurons, and the edges represent synapses. Mathematically, a graph $G = (V, E)$ consists of a set of vertices or nodes

$V = [n] := \{1, 2, \dots, n\}$ and a set of edges $E \subset$. If the graph is undirected and non-loopy, then the edge set $E = \binom{V}{2} = \binom{[n]}{2}$, where $\binom{[n]}{2}$ denotes all (unordered) pairs $\{v_i, v_j\}$ for all $v_i, v_j \in V$ and $v_i \neq v_j$. If the graph is directed and non-loopy, there are $n(n-1)$ possible edges, and we can represent them as ordered pairs. In this work, we assume the graphs are undirected, weighted, and non-loopy. The adjacency matrix A of G is the n -by- n matrix in which each entry A_{ij} denotes the edge existence between vertex i and j : $A_{ij} = 1$ if an edge is present, and $A_{ij} = 0$ if an edge is absent. The adjacency matrix is symmetric, binary and hollow, i.e., the diagonals are all zeros.

For the hermaphrodite *C.elegans* worm, the chemical connectome G_c is weighted and directed. The electrical gap junctional connectome G_g is weighted and undirected. This is consistent with an important characteristic of electrical synapses – they are bidirectional [Purves et al., 2001]. The chemical connectome G_c has 3 loops and no isolated vertices, while the electrical gap junctional connectome G_g has no loops and 26 isolated vertices. Both connectomes are sparse. The chemical connectome G_c has 2194 directed edges out of $279 \cdot 278$ possible ordered neuron pairs, resulting in a sparsity level of approximately 2.8%. The electrical gap junctional connectome G_g has 514 undirected edges out of $\binom{279}{2}$ possible unordered neuron pairs, resulting in a sparsity level of approximately 1.3%.

In our analysis, we are interested in the $279 - 26 = 253$ non-isolated neurons in the hermaphrodite *C.elegans* somatic nervous system. Each of these 253 neurons can be classified in a number of ways, including into 3 non-overlapping connectivity based types: sensory neurons (27.96%), interneurons (29.75%) and motor neurons (42.29%). Here we will work with binary, symmetric and hollow adjacency matrices of the neural connectomes throughout. We symmetrize A by $A \leftarrow A + A^T$, then binarize A by thresholding the positive entries of A to be 1 and 0 otherwise, and finally set the diagonal entries of A to be zero. Indeed, we focus on the existence of synaptic connectomes, and the occurrence of loops is low (3 loops in G_c and none in G_g) so we can ignore it.

An image of the *C.elegans* worm body is seen in Figure 1. The pair of the neural connectomes are visualized in Figure 2. In the chemical connectome G_c , the interneurons are heavily connected to the sensory neurons. The sensory neurons are connected more frequently to the motor neurons and interneurons than amongst themselves. In the electrical gap junction potential connectome G_g , the motor neurons are heavily connected to the interneurons. The sensory neurons are connected more frequently to the motor neurons and interneurons than among themselves. The connectome dataset is accessible at <http://openconnectome.me/herm-c-elegans>. Figure 3 presents the adjacency matrices of the paired *C.elegans* connectomes.

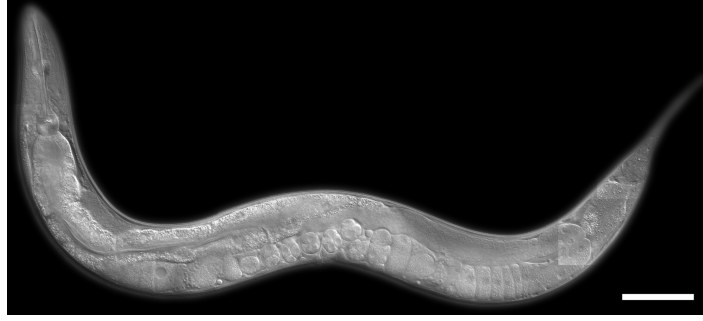


Figure 1: **An image of the *Caenorhabditis elegans* (*C.elegans*) roundworm.** The image is available at <http://post.queensu.ca/~chinsang/research/c-elegans.html>.

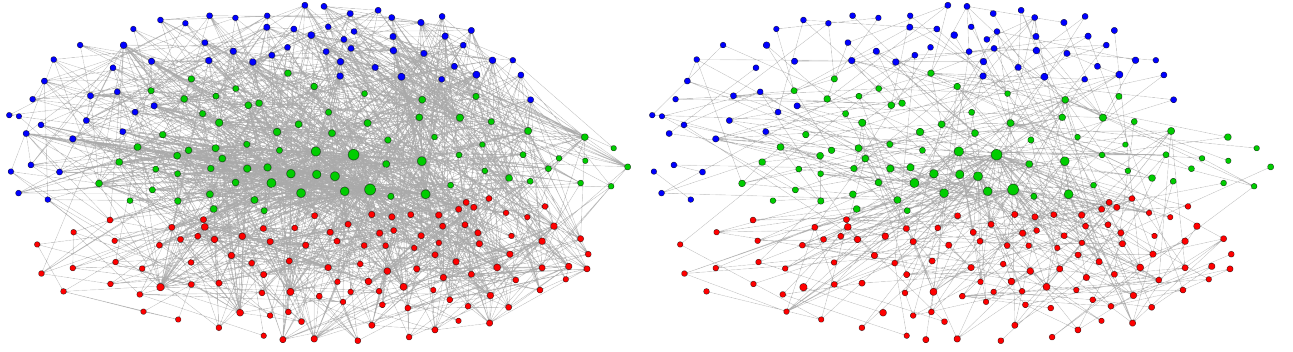


Figure 2: **The pair of *C.elegans* neural connectomes visualized as graphs.** Red nodes correspond to motor neurons, green nodes correspond to interneurons, and blue nodes correspond to sensory neurons. (Left) The chemical connectome G_c . (Right) The electrical gap junctional connectome G_g . Both synaptic connectomes are sparse, while G_g is much sparser than G_c . A similar connectivity pattern is seen across both connectomes.

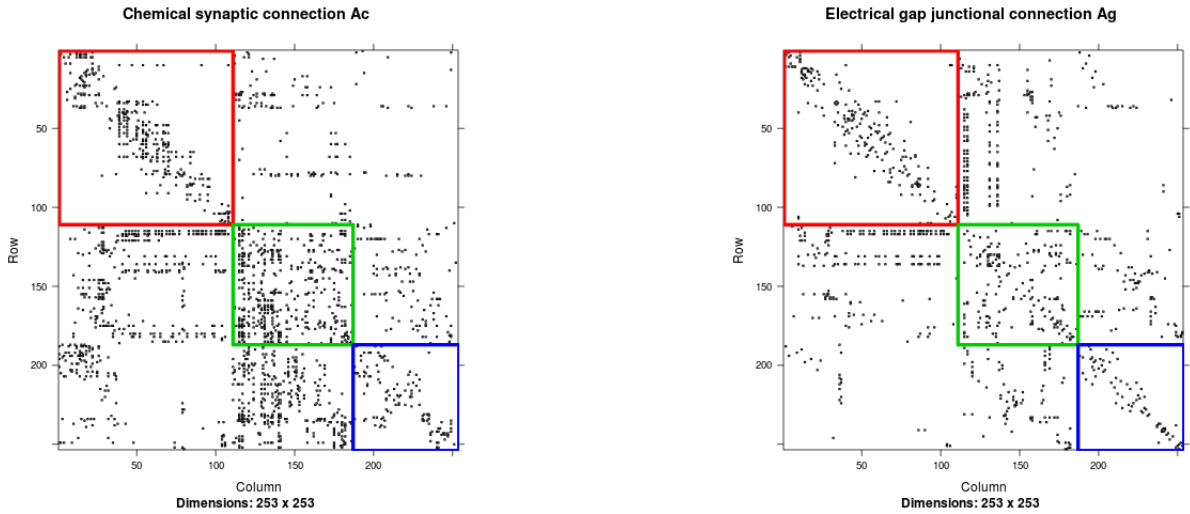


Figure 3: (Left): The adjacency matrix A_c of G_c sorted according to the neuron types. (Right): The adjacency matrix A_g of G_g sorted according to the neuron types. The red block corresponds to the connectivity among the motor neurons, the green block corresponds to the connectivity among the interneurons, and the blue block corresponds to the connectivity among the sensory neurons.

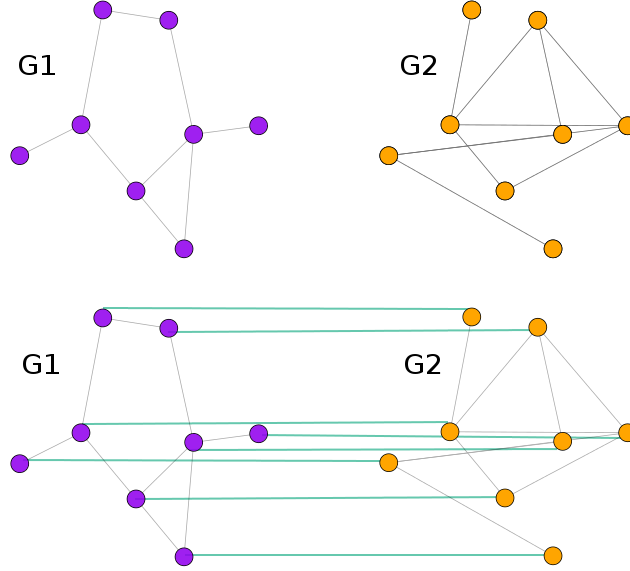


Figure 4: **A depiction of graph matching.** Given two graphs G_1 and G_2 , graph matching seeks an alignment (represented in the green lines) between the vertices across two graphs.

2 Joint Graph Inference

We consider an inference framework in the joint space of the *C.elegans* neural connectomes, which we refer to as joint graph inference. We focus on two aspects of joint graph inference: seeded graph matching and joint vertex classification.

2.1 Seeded Graph Matching

The problem of seeded graph matching (SGM) is a subproblem of the graph matching (GM) problem, which has wide applications in object recognition [Berg et al., 2005, Caelli and Kosi-
nov, 2004], image analysis [Conte et al., 2003], computer vision [Cho and Lee, 2012, Zhou and De la Torre, 2012], and neuroscience [Haris et al., 1999, Vogelstein et al., 2011, Zaslavskiy et al., 2009]. Given two graphs, $G_1 = (V_1, E_1)$ and $G_2 = (V_2, E_2)$ with respective adjacency matrices A_1 and A_2 , and $|V_1| = |V_2| = n$, the GM problem seeks a bijection ϕ between the vertex sets that minimizes edge disagreements [Lyzinski et al., 2014b, Fishkind et al., 2012]. The graph matching problem is NP-hard [Van Leeuwen and Leeuwen, 1990]. It is not known whether any graph matching algorithm is efficient, and it is suspected that none exist. For a comprehensive survey on the graph matching problem, see Conte et al. [2004] and Vogelstein et al. [2011]. The intuitive idea of graph matching is seen in Figure 4.

The seeded graph matching (SGM) problem employs additional constraint, where a partial correspondence between the vertices is known a priori. Those vertices are called “seeds”. Addition of seeds makes the graph matching problem has only a slight change in the graph

matching algorithm, and improves the graph matching performance [Fishkind et al., 2012]. Let $S_1 \subset V_1$ and $S_2 \subset V_2$ be two subsets of the vertex sets, and suppose $S_1 = S_2 = \{1, 2, \dots, m\}$. The elements of S_1 and S_2 are called seeds, and the remaining $n - m$ vertices are non-seeds. In the SGM problem, one seeks a bijection, with constraint on S_1 such that $\phi_{S_1} = \phi_{S_2}$, to minimize the number of induced edge disagreements.

The SGM problem, as a subproblem of GM, is NP-hard. We seek an approximated SGM solution that is computationally efficient. The performance of SGM solutions is measured by the matching accuracy $\delta(m)$, defined as the number of correctly matched non-seeded vertices divided by the total number of non-seeds $n - m$. When the number of seeds m is given, the remaining $n - m$ vertices need to be matched. Hence, the chance matching accuracy is $\frac{1}{n-m}$, and this accuracy increases as m increases. For larger values of m , more information on the partial correspondence between the vertices is available, and thus the SGM matching accuracy becomes higher. In this work, we apply the state-of-the-art SGM algorithm developed by Fishkind et al. [2012], seek the correspondence between the two types of neuron connectomes, and study the joint structure of the worm neural connectomes.

2.2 Joint Vertex Classification

When we observe the adjacency matrix $A \in \{0, 1\}^{n \times n}$ on n vertices and the class labels $\{Y_i\}_{i=1}^{n-1}$ associated with the first $(n - 1)$ training vertices, the task of vertex classification is to predict the label Y of the test vertex v . In this case study, the class labels are the neuron types: motor neurons, interneurons and sensory neurons. In this work, we assume the correspondence between the vertex sets across the two graphs is known. Given two graphs $G_1 = (V, E_1)$ and $G_2 = (V, E_2)$ where $V = \{v_1, \dots, v_{n-1}, v\}$, and given the class labels $\{Y_i\}_{i=1}^{n-1}$ associated with the first $(n - 1)$ training vertices, the task of joint vertex classification predicts the label of a test vertex v using information jointly from G_1 and G_2 .

Fusion inference merges information on multiple disparate data sources in order to obtain more accurate inference than using only single source. Our joint vertex classification consists of two main steps: first, a fusion information technique, namely the omnibus embedding methodology by Priebe et al. [2013]; and secondly, the inferential task of vertex classification.

The step of omnibus embedding proceeds as follows. Given G_1 and G_2 , we construct an omnibus matrix $M = \begin{pmatrix} A_1 & \Lambda \\ \Lambda & A_2 \end{pmatrix} \in \mathbb{R}^{2n \times 2n}$, where A_1 and A_2 are the adjacency matrices of G_1 and G_2 respectively, and the off-diagonal block is $\Lambda = \frac{1}{2}(A_1 + A_2)$. We consider adjacency spectral embedding [Sussman et al., 2012] of M as $2n$ points into \mathbb{R}^d . Let $U = \begin{bmatrix} U_1 \\ U_2 \end{bmatrix} \in \mathbb{R}^{2n \times d}$ denote the resulted joint embedding, where $U_1 \in \mathbb{R}^{n \times d}$ is the joint embedding corresponding to G_1 , and $U_2 \in \mathbb{R}^{n \times d}$ to G_2 . Our inference task is vertex classification. Let $\mathcal{T}_{n-1} := U_1([n - 1], :$

$) \in \mathbb{R}^{(n-1) \times d}$ denote the training set containing the first $n - 1$ vertices. We train a classifier on \mathcal{T}_{n-1} , and classify the test vertex v . A depiction of the joint vertex classification procedure is seen in Figure 5.

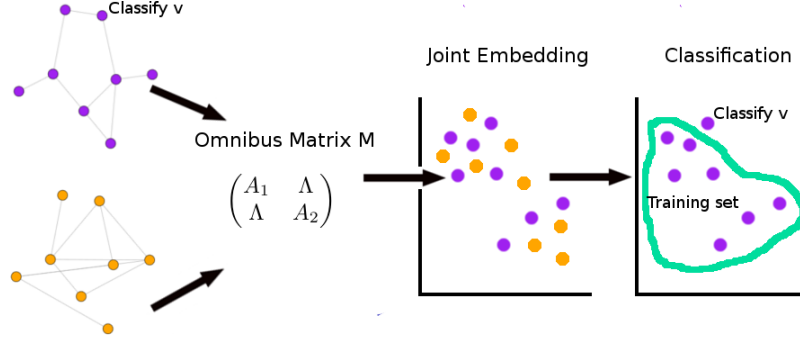


Figure 5: **A depiction of joint vertex classification.** An illustration of joint vertex classification, which embeds the joint adjacency matrix – the omnibus matrix, and classifies on the embedded space.

We demonstrate that fusing both pairs of the neural connectomes generates more accurate inference results than using a single source of connectome alone. We consider single vertex classification for comparison, which embeds the adjacency matrix A_1 to \mathbb{R}^d via adjacency spectral embedding, and classifies on the embedded space. A depiction of the single vertex classification procedure is seen in Figure 6.

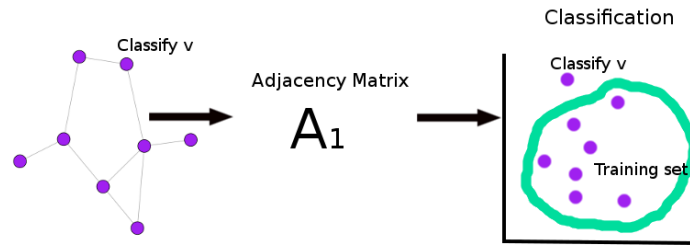


Figure 6: **A depiction of single vertex classification.** An illustration of single vertex classification, which embeds one single adjacency matrix, and classifies on the embedded space.

3 Discoveries from the Joint Space of the Neural Connectomes

3.1 Finding the Correspondence between the Chemical and the Electrical Connectomes

We apply seeded graph matching on the paired *C.elegans* neural connectomes, and discover the underlying structure preserved across the chemical and the electrical connectomes [Fishkind et al., 2012]. Figure 7 presents the errorbar plot of the seeded graph matching accuracy $\delta(m)$, plotted in black, against the number of seeds $m \in \{0, 20, 40, \dots, 180\}$. For each selected number of seeds m , we randomly and independently select 100 seeding sets S_1 . For each seeding set S_1 at a given number of seeds m , we apply the state-of-the-art seeded graph matching algorithm [Fishkind et al., 2012]. The mean accuracy $\delta(m)$ is obtained by averaging the accuracies over the 100 Monte Carlo replicates at each m . As m increases, the matching accuracy improves. This is expected, because more seeds give more information, making the SGM problem less difficult. The chance accuracy, plotted in brown dashed line, at each m is $\frac{1}{n-m}$, which does not increase significantly as m increases.

We note two significant neurological implications based on our graph matching result. First, SGM on the pair of connectomes indicate that the chemical and the electrical connectomes have statistically significant similar connectivity structure. The second significant implication is: If the performance of SGM on the chemical and the electrical connections were perfect, then one could consider just one (either one) of the paired neural connectomes without losing much information. If performance of SGM on the chemical and the electrical connections were no better than random vertex alignment, then it suggests that there is no structure similarity across the two connectomes, and this further suggests that analysis on the connectomes should proceed separately and individually. In fact, the seeded graph matching result on the *C.elegans* neural connectome is much more significant than chance but less than a perfect matching. This demonstrates that the optimal inference should be performed in the joint space of the chemical and the electrical connectomes. This discovery is noted in [Lyzinski et al., 2014a].

3.2 Predicting Neuron Types from the Joint Space of the Chemical and the Electrical Connectomes

The result of SGM on the *C.elegans* neural connectomes demonstrates the advantage of inference in the joint space of the neural connectomes, and provides a statistical motivation to apply our proposed joint vertex classification approach. Furthermore, the neurological motivation of applying joint vertex classification stems from illustrating a methodological framework

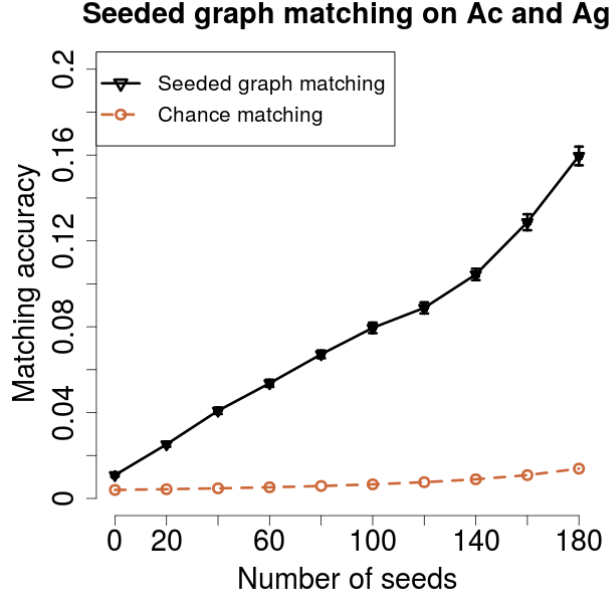


Figure 7: **Seeded graph matching on the *C.elegans* neural connectomes.** For each selected number of seeds $m \in \{0, 20, 40, \dots, 180\}$, we randomly select 100 independent seeding sets S_1 and apply SGM for each Monte Carlo replicate. The SGM mean accuracy $\delta(m)$, plotted in black, is obtained by averaging the accuracies over the 100 Monte Carlo replicates. As the number of seeds m increases, the accuracy increases. The chance accuracy, plotted in brown dashed line, is much lower than the SGM accuracy. This suggests that a significant similarity exists between the two types of synapse connections. The SGM performance on the *C.elegans* neural connectome is much more significant than chance but less than a perfect matching, indicating the optimal inference must proceed in the joint space of both neural connectomes.

to understand the coexistence and significance of chemical and electrical synaptic connectomes.

We apply joint vertex classification and single vertex classification on the paired *C.elegans* neural connectomes, and compare the classification performance. The validation is done via leave-one-out cross validation. Here we do not investigate which embedding dimension d or classifier are optimal for our classification task, and we choose support vector machine classifier with radial basis [Cortes and Vapnik, 1995] for the classification step. The paired plots in Figure 8 present the misclassification errors against the embedding dimensions $d \in \{2, 5, 8, \dots, 116, 119\}$. For the chemical connectome, the joint vertex classification (plotted in black) outperforms the single vertex classification (plotted in magenta) at all the considered embedding dimensions. For the electrical connectome, the joint vertex classification (plotted in black) outperforms the single vertex classification (plotted in magenta) at most of the considered embedding dimensions, especially larger valued embedding dimensions.

The superior performance of the joint vertex classification over the single vertex classification has an important neuroscientific implication. In many animals, the chemical synapses co-exist with the electrical synapses. Modern understanding of coexistence of chemical and

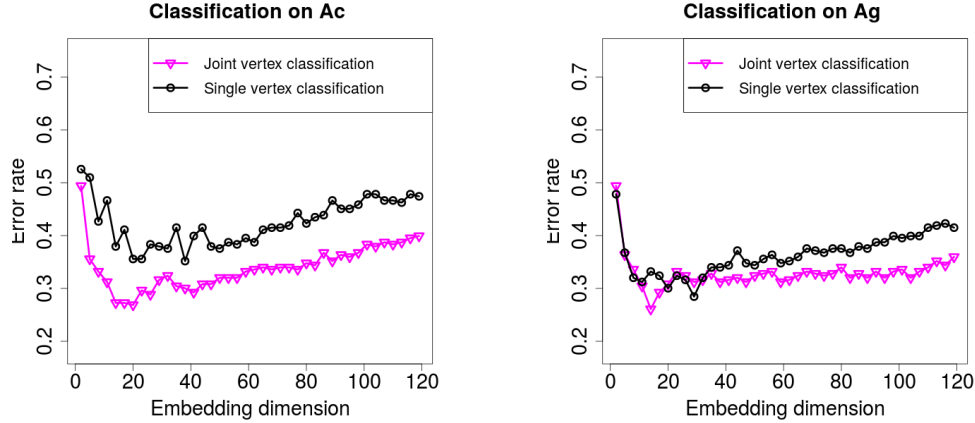


Figure 8: **Classification performance of joint and single vertex classification.** (Left) Classification on the chemical connectome A_c . For all embedding dimensions $d \in \{2, 5, 8, \dots, 116, 119\}$, the error rate of joint vertex classification, plotted in magenta, is lower than the single vertex classification, plotted in black. (Right) Classification on the electrical gap junctional connectome A_g . For most of the embedding dimensions especially with larger values, the error rate of joint vertex classification, plotted in magenta, is lower than the single vertex classification, plotted in black. Our classification result indicates that using information from the joint space of the neural connectomes improves classification performance.

electrical synaptic connectomes suggest such a coexistence has physiological significance. We discover that using both chemical and electrical connectomes jointly generates better classification performance than using one connectome alone. This may serve as a first step towards providing a methodological and quantitative approach towards understanding the coexistent significance.

4 Summary and Discussion

The paired *Caenorhabditis elegans* connectomes have become a fascinating dataset for motivating a better understanding of the nervous connectivity systems. We have presented the unique statistical approach of joint graph inference – inference in the joint graph space – to study the worm’s connectomes. Utilizing jointly the chemical and the electrical connectomes, we discover statistically significant similarity preserved across the two synaptic connectome structures. Our result of seeded graph matching indicates that the optimal inference on the information-processing properties of the connectomes must proceed in the joint space of the paired graphs.

The development of seeded graph matching provides a strong statistical motivation for joint vertex classification, where we predict neuron types in the joint space of the paired connectomes. Joint vertex classification outperforms the single vertex classification against all embedding dimensions for our different choices of dissimilarity measures. Fusion inference

using both the chemical and the electrical connectomes produces more accurate results than using one (either one) connectome alone, and enhances our understanding of the *C.elegans* connectomes. The chemical and the electrical synapses are known to coexist in most organisms. Our proposed joint vertex classification provides a methodological and quantitative framework for understanding the significance of the coexistence of the chemical and the electrical synapses. Further development of joint graph inference is a topic of ongoing investigation in both neuroscience and statistics.

5 Acknowledgement

This work is partially supported by a National Security Science and Engineering Faculty Fellowship (NSSEFF), Johns Hopkins University Human Language Technology Center of Excellence (JHU HLT COE), the XDATA program of the Defense Advanced Research Projects Agency (DARPA) administered through Air Force Research Laboratory contract FA8750-12-2-0303, and Acheson J. Duncan Fund for the Advancement of Research in Statistics.

References

- Alexander C Berg, Tamara L Berg, and Jitendra Malik. Shape matching and object recognition using low distortion correspondences. In *Computer Vision and Pattern Recognition, 2005. CVPR 2005. IEEE Computer Society Conference on*, volume 1, pages 26–33. IEEE, 2005.
- Terry Caelli and Serhiy Kosinov. An eigenspace projection clustering method for inexact graph matching. *Pattern Analysis and Machine Intelligence, IEEE Transactions on*, 26(4):515–519, 2004.
- Minsu Cho and Kyoung Mu Lee. Progressive graph matching: Making a move of graphs via probabilistic voting. In *Computer Vision and Pattern Recognition (CVPR), 2012 IEEE Conference on*, pages 398–405. IEEE, 2012.
- Donatello Conte, Pasquale Foggia, Carlo Sansone, and Mario Vento. Graph matching applications in pattern recognition and image processing. In *ICIP (2)*, pages 21–24, 2003.
- Donatello Conte, Pasquale Foggia, Carlo Sansone, and Mario Vento. Thirty years of graph matching in pattern recognition. *International journal of pattern recognition and artificial intelligence*, 18(03):265–298, 2004.
- Corinna Cortes and Vladimir Vapnik. Support-vector networks. *Machine learning*, 20(3):273–297, 1995.

- Andrew Fire, SiQun Xu, Mary K Montgomery, Steven A Kostas, Samuel E Driver, and Craig C Mello. Potent and specific genetic interference by double-stranded rna in *caenorhabditis elegans*. *nature*, 391(6669):806–811, 1998.
- Donniell E Fishkind, Sancar Adali, and Carey E Priebe. Seeded graph matching. *arXiv preprint arXiv:1209.0367*, 2012.
- Kostas Haris, Serafim N Efstratiadis, Nicos Maglaveras, Costas Pappas, John Gourassas, and George Louridas. Model-based morphological segmentation and labeling of coronary angiograms. *Medical Imaging, IEEE Transactions on*, 18(10):1003–1015, 1999.
- Rosalind C Lee, Rhonda L Feinbaum, and Victor Ambros. The *c. elegans* heterochronic gene *lin-4* encodes small rnas with antisense complementarity to *lin-14*. *cell*, 75(5):843–854, 1993.
- Vince Lyzinski, Sancar Adali, Joshua T Vogelstein, Youngser Park, and Carey E Priebe. Seeded graph matching via joint optimization of fidelity and commensurability. *arXiv preprint arXiv:1401.3813*, 2014a.
- Vince Lyzinski, Daniel Sussman, Donniell Fishkind, Henry Pao, Li Chen, Joshua T Vogelstein, Youngser Park, and Carey E Priebe. Spectral clustering for divide-and-conquer graph matching. *IEEE Parallel Computing, Systems and Applications. To appear. arXiv preprint arXiv:1310.1297*, 2014b.
- E Maupas. Modes et formes de reproduction des nematodes. *Archives de Zoologie expérimentale et générale*, 8:463–624, 1901.
- Günther Osche. Die bedeutung der osmoregulation und des winkverhaltens für freilebende nematoden. *Zeitschrift für Morphologie und ökologie der Tiere*, 41(1):54–77, 1952.
- Dragana M Pavlovic, Petra E Vértés, Edward T Bullmore, William R Schafer, and Thomas E Nichols. Stochastic blockmodeling of the modules and core of the *caenorhabditis elegans* connectome. *PloS one*, 9(7):e97584, 2014.
- Carey E Priebe, David J Marchette, Zhiliang Ma, and Sancar Adali. Manifold matching: Joint optimization of fidelity and commensurability. *Brazilian Journal of Probability and Statistics*, 27(3):377–400, 2013.
- Dale Purves, George J Augustine, David Fitzpatrick, William C Hall, Anthony-Samuel LaMantia, James O McNamara, and Leonard E White. Neuroscience. *Sunderland, MA: Sinauer Associates*, 3, 2001.

- Donald L Riddle, Thomas Blumenthal, Barbara J Meyer, James R Priess, et al. Origins of the model. 1997.
- John E Sulston, E Schierenberg, John G White, and JN Thomson. The embryonic cell lineage of the nematode *caenorhabditis elegans*. *Developmental biology*, 100(1):64–119, 1983.
- Daniel L Sussman, Minh Tang, Donniell E Fishkind, and Carey E Priebe. A consistent adjacency spectral embedding for stochastic blockmodel graphs. *Journal of the American Statistical Association*, 107(499):1119–1128, 2012.
- Emma K Towlson, Petra E Vértés, Sebastian E Ahnert, William R Schafer, and Edward T Bullmore. The rich club of the *c. elegans* neuronal connectome. *The Journal of Neuroscience*, 33(15):6380–6387, 2013.
- Jan Van Leeuwen and Jan Leeuwen. *Handbook of theoretical computer science: Algorithms and complexity*, volume 1. Elsevier, 1990.
- Lav R Varshney, Beth L Chen, Eric Paniagua, David H Hall, and Dmitri B Chklovskii. Structural properties of the *caenorhabditis elegans* neuronal network. *PLoS computational biology*, 7(2):e1001066, 2011.
- Joshua T Vogelstein, John M Conroy, Vince Lyzinski, Louis J Podrazik, Steven G Kratzer, Eric T Harley, Donniell E Fishkind, R Jacob Vogelstein, and Carey E Priebe. Fast approximate quadratic programming for large (brain) graph matching. *arXiv preprint arXiv:1112.5507*, 2011.
- Junying Yuan, Shai Shaham, Stephane Ledoux, Hilary M Ellis, and H Robert Horvitz. The *c. elegans* cell death gene *ced-3* encodes a protein similar to mammalian interleukin-1 β -converting enzyme. *Cell*, 75(4):641–652, 1993.
- Mikhail Zaslavskiy, Francis Bach, and Jean-Philippe Vert. Global alignment of protein–protein interaction networks by graph matching methods. *Bioinformatics*, 25(12):i259–1267, 2009.
- Feng Zhou and Fernando De la Torre. Factorized graph matching. In *Computer Vision and Pattern Recognition (CVPR), 2012 IEEE Conference on*, pages 127–134. IEEE, 2012.